87-130843/19 \*EP -221-844-A CIRA GEIGY AG 01.10.85-CH-004245 /13.05.87J A01n-43/40 C07d-213/30

New t-phenaxy-2-pyridyl-alkanane and-alkanal derivs. - useful os fungicides, bactericides and plant growth regulators C87-054365 EIAT BE CH DEES FR GB GR IT LI LU NL SE)

Phonoxyalkyl-pyridine derivs, of formula (1) are new:

(I)

R<sub>1</sub> - R<sub>3</sub> = H, halo, t-5C alkyl or 1-6C alkoxy (both opt. substd. by halo), CN, t-6C alkoxycarhonyl or

R, and R7 = H, 1-6C alkyl, 3-6C alkenyl, 3-6C alkynyt, or phenyl or benzyl (both opt. ring-substd. by halo, 1-6C alkyl or 1-6C alkoxy, but a opt. substd. by halo);

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Ra = H. 1-6C alkyl, 3-6C alkenyl, 3-6C alkynyl, or benzyl (opt. ring-substd. by halo, 1-6C alkyt or 1-6C alkoxy. both opt, substd. by halo); provided that the CO gp. in Rg must be in the 3- or 4-

position when  $R_1$ ,  $R_2$ ,  $R_4$ ,  $R_5$  and  $R_7$  are all H,  $R_3$  = McO and  $R_6$  = Me; and  $R_9$  can also be  $R_{10}$ CO;

R<sub>10</sub> = 1-6C alkyt (opt. substd. by halo), 3-6C alkenyl or alkynyl, 2-5C alkoxy-alkyl, 3-6C cycloalkyt (opt. substd. by 1-3C alkyl) or phenyl, benzyl or phenethyl (opt. ring-substd. by halo, 1-6C alkyl or alkoxy, both

USE/ADVANTAGE

opt. substd. by helo).

(1) are microbioldes, effective against phytopathogenic bacteria and fungi; they have curative, systemic and esp.

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preventative properties and can be applied to plants, seeds or soils. Some (I) also have plant-growth regulating activity and at higher doses inhibit excessive vegetative growth of erops.

Pref. application rates are 150-600 g/ha.

SPECIFICALLY CLAIMED

9 Cpds. c.g.

Q = H, Me, MeCO or MeO.CH, CO.

PREPARATION

Ar = phenyl substd. by R<sub>1</sub> to R<sub>3</sub>; R' = 1-4C alkyl, 3-4C alkenyl, or phenyl or benzyl, opt.

substd. by alkyl, alkoxy, halo, NO, or CN.

Reaction is pref. at -130 to 20°C, with Mg (in the form of a Grignard reagent) or Bull as metallising agent. (2)

$$\begin{array}{c|cccc}
O & R_b \\
0 & C & - \text{Hal} & \text{HO-Ar} & \frac{\text{base}}{} & \text{O}
\end{array}$$

Reaction Is pref. at 6-120°C. Both methods produce ketones which can be reduced conventionally to alcohols and these out, alkylated or acylated.

EXAMPLE

140.2 g 93% 2,4-dichlorophenyl and 232 g K<sub>2</sub>CO<sub>3</sub> were mixed in 1 l acetone, then heated briefly to boiling, cuoled to 0°C and gradualty treated over 1 hr. with 224.8% 3-(bromoacetyl)pyridine hydrobromide.

The mixt, was stirred for 15 hr. at 6-5°C and for 6 hr. at 20°C, then filtered and the mixt, evaporated. Recrystaof the residue from MeOH gave 2-(2,4-dichtorophenoxy)-1-(3-pyridinyl)-1-ethanone, m.pt. 118-9°C. (31pp1251DAHDwgNo0/0).

(G: ISR: DE2742173 EP-117485 DE2909754.

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